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GenCore version 5.1.3  
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OM protein - nucleic search, using frame\_plus\_p2n model

Run on: November 9, 2002, 07:31:31 ; Search time 299 Seconds  
(without alignments)  
1431.036 Million cell updates/sec

Title: US-09-895-298A-83  
Perfect score: 190  
Sequence: 1 MMNFQPSKMRASQMMTF.....HDGSLDRSRVSQEGNPRA 190

Scoring table:

OLIGO	
Xgapop 60.0 , Xgapext 60.0	
Ygapop 60.0 , Ygapext 60.0	
Egapop 6.0 , Egapext 7.0	
Delop 6.0 , Delext 7.0	

Searched: 2185239 segs, 1125999159 residues

Word size: 4

Total number of hits satisfying chosen parameters: 1316744

Minimum DB seq length: 0  
Maximum DB seq length: 200000000

Post-processing: Listing first 45 summaries

Command line parameters:  
-MODEL=frame+p2n.model -DEV=xlp  
-Q=/cgn2\_1/USPTO.spool/US09895298/runat\_06112002\_160752\_3557/app.query.fasta.1.327  
-DB=N\_Geneseq\_101002 -QFMT=fastq -SUFFIX=oligna.rng -MINMATCH=0.1 -LOOPCL=0  
-LOOPEXT=0 -UNITS=bits -STAR=1 -END=1 -MATRIX=oligo -TRANS=human40.cdi  
-LIST=45 -DOCALIGN=200 -THR.SCORE=quality -THR\_MIN=4 -ALIGN=15 -MODE=LOCAL  
-OUTFMT=pic -NORM-ext -HEAPSIZE=500 -MINLEN=0 -MAXLEN=200000000  
-USER=US09895298\_@CGN\_1\_1\_79\_@runat\_06112002\_160752\_3557 -NCPU=6 -ICPU=3  
-NO\_XLPXY -NO\_MAP -LARGEQUERY -NEG\_SCORES=0 -WAIT -LONGLOG -DEV.TIMEOUT=120  
-WARN\_TIMEOUT=30 -THREADS=1 -XGAPOP=60 -XGAPEXT=60 -FGAPOP=6 -FGAPEXT=7  
-YGAPOP=60 -YGAPEXT=60 -DELOP=6 -DELEXT=7

Database : N\_Geneseq\_101002:\*

1:	/SID2/gcgdata/geneseq/geneseqn-emb1/NA1980.DAT:*
2:	/SID2/gcgdata/geneseq/geneseqn-emb1/NA1981.DAT:*
3:	/SID2/gcgdata/geneseq/geneseqn-emb1/NA1982.DAT:*
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9:	/SID2/gcgdata/geneseq/geneseqn-emb1/NA1988.DAT:*
10:	/SID2/gcgdata/geneseq/geneseqn-emb1/NA1989.DAT:*
11:	/SID2/gcgdata/geneseq/geneseqn-emb1/NA1990.DAT:*
12:	/SID2/gcgdata/geneseq/geneseqn-emb1/NA1991.DAT:*
13:	/SID2/gcgdata/geneseq/geneseqn-emb1/NA1992.DAT:*
14:	/SID2/gcgdata/geneseq/geneseqn-emb1/NA1993.DAT:*
15:	/SID2/gcgdata/geneseq/geneseqn-emb1/NA1994.DAT:*
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18:	/SID2/gcgdata/geneseq/geneseqn-emb1/NA1997.DAT:*
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20:	/SID2/gcgdata/geneseq/geneseqn-emb1/NA1999.DAT:*
21:	/SID2/gcgdata/geneseq/geneseqn-emb1/NA2000.DAT:*
22:	/SID2/gcgdata/geneseq/geneseqn-emb1/NA2001A.DAT:*
23:	/SID2/gcgdata/geneseq/geneseqn-emb1/NA2001B.DAT:*
24:	/SID2/gcgdata/geneseq/geneseqn-emb1/NA2002.DAT:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description
1	190	100.0	1097	22	ABA08605	Human LAK-4p homol
2	190	100.0	1097	22	AAK53221	Human polynucleoti
3	190	100.0	1219	22	AAF82463	Human CASB6411-rel
4	190	100.0	1312	22	AAK52237	Human polynucleoti
5	190	100.0	1461	21	AAAF8402	Human secreted pro
6	190	100.0	1813	22	AAH18131	Human cDNA sequenc
7	190	100.0	1960	22	AAF82462	Human CASB6411-rel
8	190	100.0	2243	21	AAA64684	cDNA encoding a hu
9	190	100.0	2407	22	AAF82460	Human CASB6411 cDN
10	190	100.0	2521	22	AAF82461	Alternatively sp11
11	150	78.9	1194	23	ABV22463	Human prostate exp
12	150	78.9	1194	23	ABV25683	Human prostate exp
13	150	78.9	1194	23	ABV28278	Human prostate exp
14	120	63.2	470	22	AAH18491	Human breast cance
15	110	57.9	501	22	AAH09919	Human breast cance
16	71	37.4	286	23	ABV08852	Human prostate exp
17	71	37.4	617	23	ABV12915	Human prostate exp
18	52	27.4	590	23	ABV34041	Human prostate exp
19	52	27.4	590	23	ABV42908	Human prostate exp
20	50	26.3	777	22	AAH08034	Human cDNA clone (
21	46	24.2	197	22	AAH19767	Human breast cance
22	40	21.1	555	22	AAH20351	Human breast cance
23	35	18.4	454	22	ABA58847	Human foetal liver
24	35	18.4	454	22	AAK07004	Human brain expres
25	35	18.4	454	22	AAK32745	Human bone marrow
26	35	18.4	454	22	AAI38558	Probe #7244 used t
27	35	18.4	454	24	ABSO7543	Human genome-deriv
28	34	17.9	233	22	AAH10187	Human breast cance
29	32	16.8	498	22	AAH11452	Human breast cance
30	31	16.3	94	22	ABA71379	Human foetal liver
31	31	16.3	94	22	AAK19686	Human brain expres
32	31	16.3	94	22	AAK45716	Human bone marrow
33	31	16.3	94	22	AAI51641	Probe #20327 used
34	31	16.3	94	23	ABV03746	Human genome-deriv
35	29	15.3	523	23	ABV03746	Human prostate exp
36	16	8.4	402	22	AAF65737	Novel human polynu
37	16	8.4	617	22	AAH11030	Human breast cance
38	16	8.4	618	22	AAH18794	Human breast cance
39	12	6.3	775	22	AAH20104	Human breast cance
40	9	4.7	285	16	AAT25867	Human gene signatu
41	9	4.7	330	22	AAAS04641	Gene expression pr
42	9	4.7	506	24	ABQ32690	Oligonucleotide fo
43	9	4.7	506	24	ABQ32691	Oligonucleotide fo
44	9	4.7	1391	21	AACT7219	Human ORFX ORF2774
45	9	4.7	1399	22	AAH48003	Ribosomal S9 prote

ALIGNMENTS

RESULT 1  
ABAB08605  
ID ABA08605 standard; cDNA; 1097 BP.  
XX  
AC ABA08605;  
XX  
DT 11-JAN-2002 (first entry)  
DE Human LAK-4p homologue-encoding cDNA, SEQ ID NO:381.  
XX  
KW Human: cytokine; cell proliferation; cell differentiation; growth factor;  
KW haematopoiesis regulation; tissue growth; immunomodulator; activin;  
KW inhibin; chemotaxis; chemokinesis; thrombolysis; oncogenesis;  
KW proliferation; metastasis; cancer; tumour; haematopoietic disorder;  
KW myeloid cell disorder; lymphoid cell disorder; asthma; arthritis;  
KW chronic inflammatory condition; proliferative retinopathy;  
KW atherosclerosis; coronary heart disease; arterial ischemia;  
KW bone disorder; osteoporosis; vascular growth disorder;

KW tissue regeneration; wound healing; infection; immune disorder;  
KW cell culture; drug screening; gene therapy; antiinflammatory;  
KW antiasthmatic; antiarthritic; haemostatic; antiarteriosclerotic;  
KW cytosolic; osteopathic; vasotropic; cardiant; virucide; antibacterial;  
KW antifungal; vulnery; antilucer; ss.  
XX  
OS Homo sapiens.  
XX  
PN WO200157188-A2.  
XX  
PD 09-AUG-2001.  
XX  
PE 05-FEB-2001; 2001WO-US03800.  
XX  
PR 03-FEB-2000; 2000US-0496914.  
PR 27-APR-2000; 2000US-0560875.  
XX  
PA (HYSE-) HYSEQ INC.  
XX  
PI Tang YT, Liu C, Drmanac RT;  
XX  
DR WPI; 2001-457740/49.  
DR P-PSDB; ABB11361.  
XX  
PT Human proteins and DNA encoding sequences useful for preventing,  
PT treating or ameliorating a medical condition in a mammalian subject  
PT e.g. arthritis and cancer -  
XX  
PS Claim 1; Page 473; 1963pp; English.  
XX  
CC Sequences ABB10981-ABB12330 represent 1350 novel human polypeptides, and  
CC sequences ABA08225-ABA09574 represent nucleic acids encoding them. The  
CC invention also relates to vectors and recombinant host cells comprising a  
CC nucleotide of the invention, methods of producing the novel polypeptides,  
CC antibodies against the polypeptides, methods of detecting the nucleotides  
CC or polypeptides in a sample, and methods of identifying compounds which  
CC bind to polypeptides of the invention. Although novel, many of the  
CC polypeptides of the invention have homology to known proteins, thereby  
CC giving an insight into their probable biological activities, and hence  
CC potential therapeutic applications. The polypeptides of the invention may  
CC have various activities, including cytokine, cell proliferation or cell  
CC differentiation activities; stem cell growth factor activity;  
CC haematopoiesis regulatory activity; tissue growth activity;  
CC immunomodulatory activity; activin- or inhibin-related activities;  
CC chemotactic or chemokinetic activities; haemostatic, thrombotic or  
CC thrombolytic activities; receptor or ligand activities; or may be  
CC involved in oncogenesis, cancer cell proliferation or metastasis.  
CC Depending on their biological activities, polypeptides and nucleotides of  
CC the invention are useful for preventing, treating or ameliorating medical  
CC conditions, e.g., by protein or gene therapy. Such conditions include  
CC cancers, haematopoietic disorders (e.g., myeloid or lymphoid cell  
CC disorders), chronic inflammatory conditions (e.g., asthma or arthritis),  
CC proliferative retinopathy, atherosclerosis, coronary heart disease,  
CC arterial ischaemia, bone disorders (e.g., osteoporosis), and abnormal  
CC vascular growth. Polypeptides involved with tissue regeneration and  
CC repair (or nucleic acids encoding them) may be used to promote wound  
CC healing (e.g., of burns, incisions and ulcers), while those with  
CC immunomodulatory activities may be used in the treatment of viral,  
CC bacterial and fungal infections in addition to immune disorders.  
CC Polypeptides with growth factor activity may be used in cell cultures to  
CC promote cell growth. For example, such polypeptides may be used to  
CC manipulate stem cells in culture to give rise to neuroepithelial cells  
CC that can be used to augment or replace cells damaged by illness,  
CC autoimmune disease or accidental damage. The polypeptides and nucleotides  
CC may also be used in the diagnosis of the above conditions, and in drug  
CC screening techniques. The present sequence represents a cDNA encoding a  
CC novel human polypeptide of the invention.  
XX  
SQ Sequence 1097 BP; 288 A; 246 C; 247 G; 316 T; 0 other;

## Alignment Scores:

Pred. No.: 2.13e-183 Length: 1097  
Score: 190.00 Matches: 190

Percent Similarity: 100.00% Conservative: 0  
Best Local Similarity: 100.00% Mismatches: 0  
Query Match: 100.00% Indels: 0  
DB: 22 Gaps: 0

US-09-895-298A-83 (1-190) x ABA08605 (1-1097)

QY 1 MetMetAsnPhgInProProSerIysAlaTrpArgAlaSerGlnMetMetThrPhePhe 20  
DB 269 ATGATGAATTTCCAGCCTCCGAGCAAAAGCCTGGCGGCTTCACAGATGATGACTTCTTC 328  
QY 21 IlePheLeuLeuPhePheProSerPheThrGlyValLeuCysThrLeuAlaIleThrIle 40  
DB 329 ATCTTCTGCTCTTTTCCATCCTTCACCGGGGCTTGTCACACCTGGCCATCACCATC 388  
QY 41 TrpArgLeuLysProSerAlaAspCysGlyProPheArgGlyLeuPheLeuIleHis 60  
DB 389 TGGAGATTGAAGCCTTCAGCTGACTGTGGCCCTTTTGAAGGTCTGCTTCATTCAC 448  
QY 61 SerIleTySerTrpIleAspThrLeuSerThrArgProGlyTyTrpValValTrp 80  
DB 449 TCCATCTACAGCTGCGATCGACACCCCTAAGTACAGCGGCTGGCTTACCTGTGGTGG 508  
QY 81 IleTyArgAsnLeuIleGlySerValHisPhePhePheIleLeuThrLeuIleValLeu 100  
DB 509 ATCTATCGGAACCATTTGAAGTGTGCACCTTTTTCATCCTCACCCTCATTTGTGCTA 568  
QY 101 IleIleThrTyTrpLeuTyTrpGlnIleThrGlnGlyArgLysIleMetIleArgLeuLeu 120  
DB 569 ATCATCACCTATCTTTACTGGCAGATCACAGAGGGAAGGAAAGATTATGATAAGGCTGCTC 628  
QY 121 HisGlnGlnIleIleAsnGlnGlyLysAspLysMetPheLeuIleGlnLysLeuIleLys 140  
DB 629 CATGAGCAGATCATTAATGAGGGCAAGATAAATGTCCTGATGAGAAAATTGATCAAG 688  
QY 141 LeuGlnAspMetGlyLysLysAlaAsnProSerSerLeuValLeuGlnLysArgGlnVal 160  
DB 689 CTGCAGCATATGAGAAAGAACCAACCCACCTCCTGTTCTTGAAAGGAGAGAGGTG 748  
QY 161 GlnGlnGlnGlyPheLeuHisLeuGlyGlnHisAspGlySerLeuAspLeuArgSerArg 180  
DB 749 GAGCAACAAGGCTTTTTCATTTGGGGGAACATGATGCGACTGTGACTGGCATCTAGA 808  
QY 181 ArgSerValGlnGlnGlyAsnProArgAla 190  
DB 809 AGATCAGTTCAAGAAAGTAATCCAAAGGGCC 838  
RESULT 2  
AAK53221  
ID AAK53221 standard; cDNA; 1097 BP.  
XX  
AC AAK53221;  
XX  
DT 06-NOV-2001 (first entry)  
XX  
DE Human polynucleotide SEQ ID NO 2750.  
KW Human; cytokine; cell proliferation; cell differentiation; gene therapy;  
KW vaccine; peptide therapy; stem cell growth factor; haematopoiesis;  
KW tissue growth factor; immunomodulatory; cancer; leukaemia;  
KW nervous system disorder; arthritis; inflammation; ss.  
XX  
OS Homo sapiens.  
XX  
PN WO200157190-A2.  
XX  
PD 09-AUG-2001.  
XX  
PE 05-FEB-2001; 2001WO-US04098.  
XX  
PR 03-FEB-2000; 2000US-0496914.  
PR 27-APR-2000; 2000US-0560875.  
PR 20-JUN-2000; 2000US-0598075.

PR 19-JUL-2000; 2000US-0620325.  
PR 01-SEP-2000; 2000US-0654936.  
PR 15-SEP-2000; 2000US-0663561.  
PR 20-OCT-2000; 2000US-0693325.  
PR 30-NOV-2000; 2000US-0728422.  
XX  
PA (HYSE-) HYSEQ INC.  
XX  
PI Tang YT, Liu C, Drmanac RT, Asundi V, Zhou P, Xu C, Cao Y, Ma Y;  
PI Zhao QA, Wang D, Wang J, Zhang J, Ren F, Chen R, Wang ZW;  
PI Xue AJ, Yang Y, Wejhrman T, Goodrich R;  
XX  
DR WPI; 2001-476283/51.  
DR P-PSDB; AAM80088.  
XX  
PT Nucleic acids encoding polypeptides with cytokine-like activities,  
PT useful in diagnosis and gene therapy -  
XX  
PS Claim 1; Page 4962; 6221pp; English.  
XX  
CC The invention relates to polynucleotides (AAK51456-AAK53435) and the  
CC encoded polypeptides (AAM78323-AAM80302) that exhibit activity relating to  
CC cytokine, cell proliferation or cell differentiation or which may induce  
CC production of other cytokines in other cell populations. The  
CC polynucleotides and polypeptides are useful in gene therapy, vaccines or  
CC peptide therapy. The polypeptides have various cytokine-like activities,  
CC e.g. stem cell growth factor activity, haematopoiesis regulatory  
CC activity, tissue growth factor activity, immunomodulatory activity and  
CC activin/inhibin activity and may be useful in the diagnosis and/or  
CC treatment of cancer, leukaemia, nervous system disorders, arthritis and  
CC inflammation.  
CC Note: Records for SEQ ID NO 2110 (AAK52581), 2111 (AAK52582) and 3666  
CC (AAM80020) are omitted as the relevant pages from the sequence listing  
CC were missing at the time of publication.  
XX  
SQ Sequence 1097 BP; 288 A; 246 C; 247 G; 316 T; 0 other;

Alignment Scores:  
Pred. No.: 2.13e-183 Length: 1097  
Score: 190.00 Matches: 190  
Percent Similarity: 100.00% Conservative: 0  
Best Local Similarity: 100.00% Mismatches: 0  
Query Match: 100.00% Indels: 0  
DB: 22 Gaps: 0

US-09-895-298A-83 (1-190) x AAK53221 (1-1097)

QY 1 MetMetAsnPhgInProSerSerLysAlaTrpArgAlaSerGlnMetThrPhe 20  
DB 269 ATGATGAATTTCCAGCTCCGAGCAAAAGCTGCGGCTCAGATGATGACTTCTTC 328  
QY 21 IlePheLeuLeuPhePheProSerPheThrGlyValLeuCysThrLeuAlaIleThrIle 40  
DB 329 ATCTTCTTGCCTTTTCCCATCCCTTCACCGGGTCTTGTCACCCCTGGCCATCACCATC 388  
QY 41 TrpArgLeuLysProSerAlaAspCysGlyProPheArgGlyLeuProLeuPheIleHis 60  
DB 389 TGGAGATTTGAAGCCTTCACCTGACGTGTGGCCCTTTTCAGAGGTGCTCTTCATTCAC 448  
QY 61 SerIleTyrSerTrpIleAspThrLeuSerThrArgProGlyTyrLeuTrpValValTrp 80  
DB 449 TCCATCTACAGCTGGATCAGACACCTTAAGTACACGGCTGCTACCTGTGGGTTTGG 508  
QY 81 IleTyrArgAsnLeuIleGlySerValHisPhePhePheIleLeuThrLeuIleValLeu 100  
DB 509 ATCTATCGGAACCTCATTTGGAAGTGTGCACATCTTTTCATTCACCTCATTTGCTA 568  
QY 101 IleIleThrTyrLeuTyrTrpGlnIleThrGlnGlyArgLysIleMetIleArgLeuLeu 120  
DB 569 ATCATCACTAICTTACTTACGACAGATCACAGAGGGAAGATTAATGATTAAGGCTGCTC 628  
QY 121 HisGlnGlnIleIleAsnGlnGlyLysAspLysMetPheLeuIleGlnLysLeuIleLys 140  
|||||

DB 629 CATGACGAGATCATTAATGAGCGCAAGATAAATGTTCTGATAGAAAATATGATCAAG 688  
QY 141 LeuGlnAspMetGlnLysLysAlaAsnProSerSerLeuValLeuGlnArgGluVal 160  
DB 689 CTGCAGATATGAGAAAGCAAAACCCAGCTCAGTCTGTCTGGAAAGAGAGAGGTG 748  
QY 161 GlnGlnGlnGlyPheLeuHisLeuGlnGlyLysAspGlySerLeuAspLeuArgSerArg 180  
DB 749 GAGCAACAAGCCTTTTTCATTTGGGGGGAACATGATGACAGTCTTGACATCTAGTA 808  
QY 181 ArgSerValGlnGlnGlyLysProArgAla 190  
DB 809 AGATCAGTTCAAGAAAGTAATCAAGGGCC 838

RESULT 3  
AAF82463  
ID AAF82463 standard; cDNA; 1219 BP.  
XX  
AC AAF82463;  
XX  
DT 29-JUN-2001 (first entry)  
XX  
DE Human CASB6411-related cDNA #2.  
XX  
KW Human; CASB6411; vaccine; gene therapy; immunoprophylaxis;  
KW ovarian cancer; colon cancer; autoimmune disease; ss.  
XX  
OS Homo sapiens.  
XX  
FH Key Location/Qualifiers  
FT CDS 1..576  
FT /\*tag= a  
FT /partial  
FT /note= "this sequence does not contain a start codon"  
XX  
PN WO200123417-A2.  
XX  
PD 05-APR-2001.  
XX  
PF 27-SEP-2000; 2000WO-EP09500.  
XX  
PR 30-SEP-1999; 99GB-0023154.  
PR 07-JUL-2000; 2000GB-0016839.  
XX  
PA (SMIK ) SMITHKLINE BEECHAM BIOLOGICALS.  
XX  
PI Vinals De Bassols YC;  
XX  
DR WPI; 2001-316133/33.  
DR P-PSDB; AAB83082.  
XX  
PT Novel CASB6411 polypeptides useful in diagnostics, and as vaccines for  
PT prophylactic and therapeutic treatment of cancers, particularly ovarian  
PT and colon cancers, autoimmune diseases and related conditions -  
XX  
XX  
PS Claim 32; Page 66-67; 95pp; English.  
XX  
CC The present sequence is provided in a specification relating  
CC to CASB6411 polypeptides comprising a sequence having at least 70%  
CC identity to a sequence of 460 or 154 amino acids fully defined in  
CC the specification. CASB6411 polypeptides and polynucleotides are  
CC useful for treating a subject by immunoprophylaxis or therapy.  
CC The CASB6411 polypeptides are useful in diagnostics, and as  
CC vaccines for prophylactic and therapeutic treatment of cancers,  
CC particularly ovarian and colon cancers, autoimmune diseases and related  
CC conditions. CASB6411 polypeptides are also useful for the  
CC structure-based design of agonists, antagonists or inhibitors of the  
CC polypeptide.  
XX  
SQ Sequence 1219 BP; 346 A; 260 C; 275 G; 338 T; 0 other;

Alignment Scores: 2.36e-183 Length: 1219  
Pred. No.: 2.36e-183

Score: 190.00 Matches: 190  
Percent Similarity: 100.00% Conservative: 0  
Best Local Similarity: 100.00% Mismatches: 0  
Query Match: 100.00% Indels: 0  
DB: 22 Gaps: 0

US-09-895-298A-83 (1-190) x AAF82463 (1-1219)

QY 1 MetMetasnpheginProProserLysAlaTrpArgAlaSerGlnMetMetThrPhephe 20  
|||||  
Db 4 ATGATGAATTTCCAGCCTCCGAGCAAGCCCTGGCGGCTCAGAGATGATGACTTCTTC 63  
QY 21 IlePheLeuLeuPhePheProSerPheThrGlyValLeuCysThrLeuAlaIleThrIle 40  
|||||  
Db 64 ATCTTCTGCTCTTTTCCCATCTTTCACCGGGGCTTGTGACCCCTGGCCATCACCATC 123  
QY 41 TrpArgLeuLysProSerAlaAspCysGlyProPheArgGlyLeuProLeuPheIleHis 60  
|||||  
Db 124 TGGAGATTGAAGCCTTCAGCTGAGCTGGCCCTTTTCGAGTCTGCTCTTCATTCAC 183  
QY 61 SerIleTyrrSerTrpIleAspThrLeuSerThrArgProGlyTyrrLeuTrpValTrp 80  
|||||  
Db 184 TCCATCTACAGCTGGATCGACACCCCTAAGTACACGGCCTGACTGTGGGTTGTTGG 243  
QY 81 IleTyrrArgAsnLeuIleGlySerValHisPhePhePheIleLeuThrLeuIleValLeu 100  
|||||  
Db 244 ATCTATCGGAACCTCATTTGGAAGTGTGCACCTCTTTTCATCCTCACCCTCATTTGTGCTG 303  
QY 101 IleIleThrTyrrLeuTyrrTrpGlnIleThrGlnGlyArgGlyIleMetIleArgLeuLeu 120  
|||||  
Db 304 ATCATCACCTATCTTTACTTGGCAGATCACAGAGGGAAGGAAATTAATGATTAAGGCTGCTC 363  
QY 121 HisGlnGlnIleIleAsnGlnGlyLysAspLysMetPheLeuIleGlnLysLeuIleLys 140  
|||||  
Db 364 CATGAGCAGATCATTAATGAGGGGCAAAATGTTCTCTGATAGAAAAATGATCAG 423  
QY 141 LeuGlnAspMetGlnLysLysAlaAsnProSerSerLeuValLeuGlnArgArgGlnVal 160  
|||||  
Db 424 CTGCAGATATGGAGAAAGCAAAACCCACGCTCAGCTTGTGAAAAGGAGAGAGTG 483  
QY 161 GlnGlnGlnGlyPheLeuHisLeuGlyGlnHisAspGlySerLeuAspLeuArgSerArg 180  
|||||  
Db 484 GAGCAACAAGGCTTTTGCATTTGGGGGAACATGATGGCAGTCTTGACTTGGGATCTAGA 543  
QY 181 ArgSerValGlnGlnGlyAsnProArgAla 190  
|||||  
Db 544 AGATCAGTTCAAGAGGTAATCCAAAGGGCC 573

RESULT 4  
AAK52237  
ID AAK52237 standard; cDNA; 1312 BP.

XX AAK52237;  
AC AAK52237;  
DT 06-NOV-2001 (first entry)  
XX Human polynucleotide SEQ ID NO 782.  
DE Human; cytokine; cell proliferation; cell differentiation; gene therapy;  
KW vaccine; peptide therapy; stem cell growth factor; haematopoiesis;  
KW tissue growth factor; immunomodulatory; cancer; leukaemia;  
KW nervous system disorder; arthritis; inflammation; ss.  
XX Homo sapiens.  
XX WO200157190-A2.  
XX 09-AUG-2001.  
XX 05-FEB-2001; 2001MO-US04098.  
XX 03-FEB-2000; 2000US-0496914.  
XX 27-APR-2000; 2000US-0560875.

PR 20-JUN-2000; 2000US-0598075.  
PR 19-JUL-2000; 2000US-0620325.  
PR 01-SEP-2000; 2000US-0654936.  
PR 15-SEP-2000; 2000US-0663561.  
PR 20-OCT-2000; 2000US-0693325.  
PR 30-NOV-2000; 2000US-0728422.

XX (HYSE-) HYSEQ INC.

XX Tang YT, Liu C, Drmanac RT, Asundi V, Zhou P, Xu C, Cao Y, Ma Y;  
PI Zhao Qa, Wang D, Wang J, Zhang J, Ren F, Chen R, Wang ZW;  
PI Xue AJ, Yang Y, Wejhrman T, Goodrich R;

XX MPI: 2001-476283/51.  
DR P-PSDB; AAM79104.

XX Nucleic acids encoding polypeptides with cytokine-like activities,  
PT useful in diagnosis and gene therapy -  
PS Claim 1; Page 2615-2616; 6221pp; English.

XX The invention relates to polynucleotides (AAK51456-AAK53435) and the  
CC encoded polypeptides (AAM78323-AAM80302) that exhibit activity elating to  
CC cytokine, cell proliferation or cell differentiation or which may induce  
CC production of other cytokines in other cell populations. The  
CC polynucleotides and polypeptides are useful in gene therapy, vaccines or  
CC peptide therapy. The polypeptides have various cytokine-like activities,  
CC e.g. stem cell growth factor activity, haematopoiesis regulating  
CC activity, tissue growth factor activity, immunomodulatory activity and  
CC activin/inhibin activity and may be useful in the diagnosis and/or  
CC treatment of cancer, leukaemia, nervous system disorders, arthritis and  
CC inflammation.  
CC Note: Records for SEQ ID NO 2110 (AAK52581), 2111 (AAK52582) and 3666  
CC (AAM80020) are omitted as the relevant pages from the sequence listing  
CC were missing at the time of publication.

XX Sequence 1312 BP; 370 A; 286 C; 287 G; 369 T; 0 other;

XX Alignment Scores:

Pred. No.: 2,53e-183 Length: 1312  
Score: 190.00 Matches: 190  
Percent Similarity: 100.00% Conservative: 0  
Best Local Similarity: 100.00% Mismatches: 0  
Query Match: 100.00% Indels: 0  
DB: 22 Gaps: 0

US-09-895-298A-83 (1-190) x AAK52237 (1-1312)

QY 1 MetMetasnpheginProProserLysAlaTrpArgAlaSerGlnMetMetThrPhephe 20  
|||||  
Db 294 ATGATGAATTTCCAGCCTCCGAGCAAGCCCTGGCGGCTCAGAGATGATGACTTCTTC 353  
QY 21 IlePheLeuLeuPhePheProSerPheThrGlyValLeuCysThrLeuAlaIleThrIle 40  
|||||  
Db 354 ATCTTCTGCTCTTTTCCCATCCTTCACCGGGGCTTGTGACCCCTGGCATCACCATC 413  
QY 41 TrpArgLeuLysProSerAlaAspCysGlyProPheArgGlyLeuProLeuPheIleHis 60  
|||||  
Db 414 TGGAGATTGAAGCCTTCAGCTGAGCTGGCCCTTTTCGAGGTCGCTCTTCATTCAC 473  
QY 61 SerIleTyrrSerTrpIleAspThrLeuSerThrArgProGlyTyrrLeuTrpValTrp 80  
|||||  
Db 474 TCCATCTACAGCTGGATCGACACCCCTAAGTACACGGCCTGCTACCTGTGGTGTGG 533  
QY 81 IleTyrrArgAsnLeuIleGlySerValHisPhePhePheIleLeuThrLeuIleValLeu 100  
|||||  
Db 534 ATCTATCGGAACCTCATTTGGAAGTGTGCACCTCTTTTCATCCTCACCCTCATTTGTGCTA 593  
QY 101 IleIleThrTyrrLeuTyrrTrpGlnIleThrGlnGlyArgGlyIleMetIleArgLeuLeu 120  
|||||  
Db 594 ATCATCACCTATCTTTACTTGGCAGATCACAGAGGGAAGGAAATTAATGATTAAGGCTGCTC 653  
QY 121 HisGlnGlnIleIleAsnGlnGlyLysAspLysMetPheLeuIleGlnLysLeuIleLys 140

```
|||||
Db 654 CATGACGACATCATTAATGAGCGCAAAAGATAAATGTTCTCGATAGAAAATTGATCAAG 713
QY 141 LeuGlnAspMetGluLysAlaAsnProSerSerLeuValLeuGlnArgGluVal 160
Db 714 CTGACGATATGAGAGAAAGCAAAACCCAGCTCACTTGTCTGAAAGAGAGAGGTG 773
QY 161 GluGlnGlyPheLeuHisLeuGlyGlnHisAspGlySerLeuAspLeuArgSerArg 180
Db 774 GAGCAACAAGGCTTTTGGCATTTGGGGGACATGATGGCAGTCTTGACTTGGATCTAGA 833
QY 181 ArgSerValGlnGluGlyAsnProArgAla 190
Db 834 AGATCAGTTCAGAAGGTAAATCCAAGGCC 863

RESULT 5
AAA78402
ID AAA78402 standard; cDNA; 1461 BP.
XX
AC AAA78402;
XX
DT 20-NOV-2000 (first entry)
XX
DE Human secreted protein gene 22 SEQ ID NO:32.
XX
KW Human; secreted protein; cytostatic; antianaemic; antidiabetic;
KW antinflammatory; ophthalmological; antirheumatic; antiarthritic;
KW antipsoriatic; antiangiogenic; cardiant; anti-HIV; nootropic;
KW neuroprotective; antimicrobial; antiparkinsonian; cancer;
KW immune system disorder; angiogenesis; hyperproliferative disorder;
KW cardiovascular disorder; apoptosis; neurological disease;
KW infectious disease; wound healing; ss.
XX
OS Homo sapiens.
XX
PN WO200035937-A1.
XX
PD 22-JUN-2000.
XX
PF 16-DEC-1999; 99WO-US29950.
XX
PR 17-DEC-1998; 98US-0112809.
PR 18-DEC-1998; 98US-0113006.
XX
PA (HUMA-) HUMAN GENOME SCI INC.
XX
PI Ruben SM, Ebner R, Rosen CA, Endress GA, Soppet DR, Ni J,
PI Duan DR, Moore PA, Shi Y, Lafleur DW, Olsen HS, Florence K;
XX
DR WPI: 2000-431566/37.
DR P-PSDB; AAB24458.
XX
PT Forty seven human nucleic acids encoding secreted proteins, useful in
PT the treatment, prevention and diagnosis of cancers, disorders of the
PT immune system, angiogenesis disorders, neurological diseases and
PT hyperproliferative disorders -
XX
PS Claim 1; Page 457-458; 562pp; English.
XX
CC The polynucleotide sequence given in AAA78381 to AAA78432 encode the
CC human secreted proteins given in AAB24437 to AAB24604. Human secreted
CC proteins have activities based on the tissues and cells the genes are
CC expressed in. Examples of activities include: cytostatic; antianaemic;
CC antidiabetic; antinflammatory; ophthalmological; antirheumatic;
CC antiarthritic; antipsoriatic; antiangiogenic; cardiant; anti-HIV;
CC nootropic; neuroprotective; antimicrobial and antiparkinsonian.
CC Human secreted protein polynucleotides, polypeptides, antagonists and/or
CC agonists may be useful in treating, preventing, and/or diagnosing other
CC diseases, disorders, and/or conditions such as: (a) cancers; (b)
CC disorders of the immune system; (c) angiogenesis disorders; (d)
CC hyperproliferative disorders; (e) cardiovascular disorders; (f) diseases
CC associated with increase apoptosis; (g) neurological diseases; and
CC (h) infectious diseases. They are also used to promote wound healing.
```

```
CC AAA78372 to AAA78380 and AAB24436 represent sequences used in the
CC exemplification of the present invention.
XX
SQ Sequence 1461 BP; 428 A; 312 C; 324 G; 397 T; 0 other;

Alignment Scores:
Pred. No.: 2.81e-183 Length: 1461
Score: 190.00 Matches: 190
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 21 Gaps: 0

US-09-895-298a-83 (1-190) x AAA78402 (1-1461)

QY 1 MetMetAsnPhenGlnProProSerLysAlaTrpArgAlaSerGlnMetMetThrPhePhe 20
Db 63 ATGATGAATTTCACGCGCTCCGAGCAAAAGCCCTGGCGGCTCACAGATGATGACTTCTTC 122
QY 21 IlePheLeuLeuPhePheProSerPheThrGlyValLeuCysThrLeuAlaIleThrIle 40
Db 123 ATCTTCTTGGCTCTTTTCCCATCTTTCACCGGGGTCTTGTGCACCCCTGGCCATCACCATC 182
QY 41 TrpArgLeuLysProSerAlaAspCysGlyProPheArgGlyLeuProLeuPheIleHis 60
Db 183 TGGAGATTGAAGCCTTCAGCTGACGTGGCCCTTTCGAGAGTCTGCCCTCTTCATTCCAC 242
QY 61 SerIleTyrSerTrpIleAspThrIleSerThrArgProGlyTyrLeuTrpValIleTrp 80
Db 243 TCCATCTACAGCTGGATGACACACCCCTAAGTACACGGCTGCTACCTGTGGTGTGG 302
QY 81 IleTyrArgAsnLeuIleGlySerValHisPhePheIleLeuThrIleValLeu 100
Db 303 ATCTATCGGAACCTCATGTGAAGTGTGACCTTCTTTCATCCTCACCCCTCATTTGTGCTA 362
QY 101 IleIleThrTyrLeuTyrTrpGlnIleThrGlnGlyArgLysIleMetIleArgLeuLeu 120
Db 363 ATCATCACCTTACTTACTGGCAGATCACAGAGGAAGAAAGATATGATGAAGCGTCTC 422
QY 121 HisGlnGlnIleIleAsnGlnGlyLysAspLysMetPheLeuIleGluLysLeuIlys 140
Db 423 CATGACGACATCATTAATGAGGGCCAAAGATATAATGTTCTGTAGAAAATTGATCAAG 482
QY 141 LeuGlnAspMetGluLysAlaAsnProSerSerLeuValLeuGlnArgGluVal 160
Db 483 CTGACGATATGAGAGAAAGCAAAACCCAGCTCACTTGTCTGAAAGAGAGAGGTG 542
QY 161 GluGlnGlnGlyPheLeuHisLeuGlyGlnHisAspGlySerLeuAspLeuArgSerArg 180
Db 543 GAGCAACAAGGCTTTTGGCATTTGGGGGAAACATGATGGCAGTCTTGACTTGGATCTAGA 602
QY 181 ArgSerValGlnGluGlyAsnProArgAla 190
Db 603 AGATCAGTTCAGAAGGTAAATCCAAGGCC 632

RESULT 6
AAH18131
ID AAH18131 standard; cDNA; 1813 BP.
XX
AC AAH18131;
XX
DT 26-JUN-2001 (first entry)
XX
DE Human cDNA sequence SEQ ID NO:18001.
XX
KW Human; primer; detection; diagnosis; antisense therapy; gene therapy; ss.
XX
OS Homo sapiens.
XX
PN EP1074617-A2.
XX
PD 07-FEB-2001.
XX
```



PF 28-JUL-2000; 2000EP-0116126.  
XX  
PR 29-JUL-1999; 99JP-0248036.  
PR 27-AUG-1999; 99JP-0300253.  
PR 11-JAN-2000; 2000JP-0118776.  
PR 02-MAY-2000; 2000JP-0183767.  
PR 09-JUN-2000; 2000JP-0241899.  
XX  
PA (HELI-) HELIX RES INST.  
XX  
PI Ota T, Isogal T, Nishikawa T, Hayashi K, Saito K, Yamamoto J;  
PI Ishii S, Sugiyama T, Wakamatsu A, Nagai K, Otsuki T;  
XX  
DR WPI; 2001-318749/34.  
XX  
PT Primer sets for synthesizing polynucleotides, particularly the 5602  
PT full-length cDNAs defined in the specification, and for the detection  
PT and/or diagnosis of the abnormality of the proteins encoded by the  
PT full-length cDNAs -  
XX  
PS Claim 8; SEQ ID 18001; 2537bp + CD ROM; English.  
XX  
CC The present invention describes primer sets for synthesizing 5602  
CC full-length cDNAs defined in the specification. Where a primer set  
CC comprises: (a) an oligo-dT primer and an oligonucleotide complementary  
CC to the complementary strand of a polynucleotide which comprises one of  
CC the 5602 nucleotide sequences defined in the specification, where the  
CC oligonucleotide comprises at least 15 nucleotides; or (b) a combination  
CC of an oligonucleotide comprising a sequence complementary to the  
CC complementary strand of a polynucleotide which comprises a 5'-end  
CC sequence and an oligonucleotide comprising a sequence complementary to a  
CC polynucleotide which comprises a 3'-end sequence, where the  
CC oligonucleotide comprises at least 15 nucleotides and the combination of  
CC the 5'-end sequence/3'-end sequence is selected from those defined in  
CC the specification. The primer sets can be used in antisense therapy and  
CC in gene therapy. The primers are useful for synthesizing polynucleotides,  
CC particularly full-length cDNAs. The primers are also useful for the  
CC detection and/or diagnosis of the abnormality of the proteins encoded by  
CC the full-length cDNAs. The primers allow obtaining of the full-length  
CC cDNAs easily without any specialised methods. AAH03166 to AAH13628 and  
CC AAH13633 to AAH18742 represent human cDNA sequences; AAB92446 to  
CC AAB95893 represent human amino acid sequences; and AAH13629 to AAH13632  
CC represent oligonucleotides, all of which are used in the exemplification  
CC of the present invention.  
XX  
SQ Sequence 1813 BP; 489 A; 400 C; 405 G; 519 T; 0 other;  
  
Alignment Scores:  
Pred. No.: 3,46e-183 Length: 1813  
Score: 190.00 Matches: 190  
Percent Similarity: 100.00% Conservative: 0  
Best Local Similarity: 100.00% Mismatches: 0  
Query Match: 100.00% Indels: 0  
DB: 22 Gaps: 0  
  
US-09-895-298A-83 (1-190) x AAH18131 (1-1813)  
QY 1 MetMetasnpheglnproproserlysalatrpargalaserglmethrphphe 20  
DB 451 ATGATGAATTCAGCCTCCAGCAAGCCTGGCGGCTCAGATGATGACTTCTTC 510  
QY 21 IlePheLeuLeuPhePheProSerPheThrglyValLeuCysThrLeuAlaIleThrIle 40  
DB 511 ATCTTCTTGCTCTTTTCCATCCTTCACCGGGGCTTGTCACACCTGGCCATCACCATC 570  
QY 41 TTPArgLeuLysProSerAlaAspCysGlyProPheArgGlyLeuProLeuPheIleHis 60  
DB 571 TGGAGATGAAGCCTTCAGCTGAGTGGGCCCTTTTCGAGGTCTGCTTCATTCAC 630  
QY 61 SerIleTySerTrpIleAspThrLeuSerThrArgProGlyTyrLeuTrpValValTrp 80  
DB 631 TTCATCTACAGCTGATGCACACCTAAGTACACGGCGCTGCTGCTGCTGCTTGTGG 690

QY 81 IleTyArgAsnLeuIleGlySerValHisPhePheIleLeuThrLeuIleValLeu 100  
DB 691 ATCTATCGGAACCTCATTTGGAAGTGTGCACCTCTTTTCATCTCCACCTCATTTGTGCTA 750  
QY 101 IleIleThrTyrrLeuTyrrTrpGlnIleThrGlyArgLysIleMetIleArgLeuLeu 120  
DB 751 ATCATCACCCTATCTTTACTGTGCAGATCAGACGAGGAAGATATGATAGGCTGCTC 810  
QY 121 HisGlnGlnIleIleAsnGlnGlyLysAspLysMetPheLeuIleGlnLysLeuIleLys 140  
DB 811 CATGAGCAGATCATTAATGAGGGCAAGATAAATGTTCTGTATGAAAAATTGATCAAG 870  
QY 141 LeuGlnAspMetGlnLysLysAlaAsnProSerSerLeuValLeuGluValArgGluVal 160  
DB 871 CTGCAGGATATGGAGAAGAAGCAACCCAGCTCAGCTGTTCTGGAAGAGAGAGAGTG 930  
QY 161 GlnGlnGlnGlyPheLeuHisLeuGlyGluHisAspGlySerLeuAspLeuArgSerArg 180  
DB 931 GAGCAACAAGCGCTTTTGCATTTGGGGGAACATGATGCAGCTTGACCTGCATCTAGA 990  
QY 181 ArgSerValGlnGlnGlyAsnProArgAla 190  
DB 991 AGATCAGTTCAAGAAGTAATCCAAGGGCC 1020  
  
RESULT 7  
AAF82462  
ID AAF82462 standard; cDNA; 1960 BP.  
XX  
AC AAF82462;  
XX  
DT 29-JUN-2001 (first entry)  
XX  
DE Human CASB6411-related cDNA #1.  
XX  
KW Human; CASB6411; vaccine; gene therapy; immunoprophylaxis;  
KW ovarian cancer; colon cancer; autoimmune disease; ss.  
XX  
OS Homo sapiens.  
XX  
FH Key Location/Qualifiers  
FT CDS 1..1317  
FT /\*tag= a  
FT /partial  
FT /note= "this sequence does not contain a start codon"  
XX  
PN WO200123417-A2.  
XX  
PD 05-APR-2001.  
XX  
PE 27-SEP-2000; 2000WO-EP09500.  
XX  
PR 30-SEP-1999; 99GB-0023154.  
PR 07-JUL-2000; 2000GB-0016839.  
XX  
PA (SMIK ) SMITHKLINE BEECHAM BIOLOGICALS.  
XX  
PI Vinals De Bassols YC;  
PI WPI; 2001-316133/33.  
PI P-PSDB; AAB83081.  
XX  
PT Novel CASB6411 polypeptides useful in diagnostics, and as vaccines for  
PT prophylactic and therapeutic treatment of cancers, particularly ovarian  
PT and colon cancers, autoimmune diseases and related conditions -  
XX  
PS Claim 32; Page 65-66; 95pp; English.  
XX  
CC The present sequence is provided in a specification relating  
CC to CASB6411 polypeptides comprising a sequence having at least 70%  
CC identity to a sequence of 460 or 154 amino acids fully defined in  
CC the specification. CASB6411 polypeptides and polynucleotides are  
CC useful for treating a subject by immunoprophylaxis or therapy.  
CC The CASB6411 polypeptides are useful in diagnostics, and as





Db 760 ATCTTCTGCTCTTTTCCATCCTTCACCGGGGCTTGTGTGCACCCCTGGCCATCACCATC 819  
QY 41 TTPARGLEULYSPROSERALIAASPCYSGLYPROPHARGGLYLEUPROLEUPHEILEHIS 60  
Db 820 TGGAGATGGAAGCCTTCAGCTGACTGTGGCCCTTTTCGAGGTCTGCCTCTTCATTCAC 879  
QY 61 SerIleTyrSerTrpIleAspThrIleuSerThrArgProGlyTyrIleuTrpValValTrp 80  
Db 880 TCCATCTACAGCTGGATGCACACCCCTAAAGTACACGGCTGGCTACTGTGGGTGTGG 939  
QY 81 IleTyrArgAsnIleuIleGlySerValHisPhePheIleuThrIleuValIleu 100  
Db 940 ATCTATCGGAACCTCATTTGGAAGTGTGCACCTTTTTCATCCTCACCCTCATTTGTGCTA 999  
QY 101 IleIleThrTyrIleuTyrTrpGlnIleThrGlnGlyArgGlyIleMetIleArgIleu 120  
Db 1000 ATCATCACTTACTTTACTGCGACATCACAGAGGGAAGATATGATTAAGCTGCTC 1059  
QY 121 HisGlnGlnIleIleAsnGlnGlyLysAspLysMetPheIleuIleGlnLysIleLys 140  
Db 1060 CATGACGAGATCATTAATGAGGGCAAGATAAATGTTCTGATAGAAAAATTGATCAAG 1119  
QY 141 LeuGlnAspMetGlnLysLysAlaAsnProSerSerIleuValIleuGlnArgGlnVal 160  
Db 1120 CTGCAGGATATGAGAAGAAAGCAACCCACGCTCAGCTGTTCTCGAAAGAGAGAGGTG 1179  
QY 161 GlnGlnGlnGlyPheLeuHisLeuGlyGlnHisAspGlySerIleuAspLeuArgSerArg 180  
Db 1180 GAGCAACAGAGCTTTTTCATTTGGGGAAACATGATGGCAGCTTGACTTGCGATCTAGA 1239  
QY 181 ArgSerValGlnGlnGlyAsnProArgAla 190  
Db 1240 AGATCAGTTCAAGAAGTATCCCAAGGGCC 1269  
RESULT 9  
AAF82460  
ID AAF82460 standard; cDNA; 2407 BP.  
XX  
AC AAF82460;  
XX  
DT 29-JUN-2001 (first entry)  
XX  
DE Human CASB6411 cDNA.  
XX  
KM Human; CASB6411; vaccine; gene therapy; immunoprophylaxis;  
XX  
KM ovarian cancer; colon cancer; autoimmune disease; ss.  
XX  
OS Homo sapiens.  
XX  
FH Key Location/Qualifiers  
FT CDS 382..1764  
FT /\*tag= a  
FT /product= "CASB6411"  
XX  
XX WO200123417-A2.  
XX  
XX PD 05-APR-2001.  
XX  
XX PF 27-SEP-2000; 2000WO-EP09500.  
XX  
XX PR 30-SEP-1999; 99GB-0023154.  
XX PR 07-JUL-2000; 2000GB-0016839.  
XX  
XX PA (SMIK ) SMITHKLINE BEECHAM BIOLOGICALS.  
XX  
XX PI Vinals De Bassols YC;  
XX  
XX DR WPI; 2001-316133/33.  
XX DR P-PSDB; AAB83079.  
XX  
PT Novel CASB6411 polypeptides useful in diagnostics, and as vaccines for  
PT prophylactic and therapeutic treatment of cancers, particularly ovarian  
PT and colon cancers, autoimmune diseases and related conditions -

XX  
PS Claim 11; Page 63-64; 95pp; English.  
XX  
CC The present sequence encodes human CASB6411 polypeptide. The  
CC invention relates to CASB6411 polypeptides comprising a sequence  
CC having at least 70% identity to a sequence of 460 or 154 amino acids  
CC fully defined in the specification. CASB6411 polypeptides and  
CC polynucleotides are useful for treating a subject by immunoprophylaxis  
CC or therapy. The CASB6411 polypeptides are useful in diagnostics, and  
CC as vaccines for prophylactic and therapeutic treatment of cancers,  
CC particularly ovarian and colon cancers, autoimmune diseases and related  
CC conditions. CASB6411 polypeptides are also useful for the  
CC structure-based design of agonists, antagonists or inhibitors of the  
CC polypeptide. The present sequence may be alternatively spliced to  
CC generate a sequence encoding a truncated CASB6411 protein.  
XX  
SQ Sequence 2407 BP; 635 A; 557 C; 546 G; 669 T; 0 other;  
Alignment Scores:  
Pred. No.: 4.55e-183 Length: 2407  
Score: 190.00 Matches: 190  
Percent Similarity: 100.00% Conservative: 0  
Best Local Similarity: 100.00% Mismatches: 0  
Query Match: 100.00% Indels: 0  
DB: 22 Gaps: 0  
US-09-895-298A-83 (1-190) x AAF82460 (1-2407)  
QY 1 MetMetAsnPheGlnProProSerLysAlaTrpArgAlaSerGlnMetMetThrPhe 20  
Db 1192 ATGATGAATTTCCAGCCTCCGAGCAAGCCTGGCGGCTCAGATGATGACTTCTTC 1251  
QY 21 IlePheLeuLeuPhePheProSerPheThrGlyValIleuCysThrIleuAlaIleThrIle 40  
Db 1252 ATCTCTTGCTCTTTTCCCATCCTTCACCGGGGTCTGTGCACCTGGCCATCACCATC 1311  
QY 41 TTPARGLEULYSPROSERALIAASPCYSGLYPROPHARGGLYLEUPROLEUPHEILEHIS 60  
Db 1312 TGGAGATTGAAGCCTTCAGCTGACTGTGGCCCTTTTCGAGGTCTGCTCTTCATTCAC 1371  
QY 61 SerIleTyrSerTrpIleAspThrIleuSerThrArgProGlyTyrIleuTrpValValTrp 80  
Db 1372 TCCATCTACAGCTGGATCGACACCCCTAAGTACACGGCTGGCTACTGTGGGTGTGG 1431  
QY 81 IleTyrArgAsnIleuIleGlySerValHisPhePheIleuThrIleuIleValIleu 100  
Db 1432 ATCTATCGGAACCTCATTTGGAAGTGTGCACCTTTTTCATCCTCACCCTCATTTGTGCTA 1491  
QY 101 IleIleThrTyrIleuTyrTrpGlnIleThrGlnGlyArgGlyIleMetIleArgIleu 120  
Db 1492 ATCATCACTTATCTTTACTGCGAGATCACAGAGGGAAGATATGATTAAGCTGTGCTC 1551  
QY 121 HisGlnGlnIleIleAsnGlnGlyLysAspLysMetPheIleuIleGlnLysIleLys 140  
Db 1552 CATGACGAGATCATTAATGAGGGCAAGATAAATGTTCTGATAGAAAAATTGATCAAG 1611  
QY 141 LeuGlnAspMetGlnLysLysAlaAsnProSerSerIleuValIleuGlnArgArgGlnVal 160  
Db 1612 CTGCAGGATATGAGAAGAAAGCAACCCACGCTCAGCTTGTTCGAAAGAGAGAGGTG 1671  
QY 161 GlnGlnGlnGlyPheLeuHisLeuGlyGlnHisAspGlySerIleuAspLeuArgSerArg 180  
Db 1672 GAGCAACAAGCCTTTTTCGACTTTGGGGGAACATGATGCAGCTTGACTTGCATCTAGA 1731  
QY 181 ArgSerValGlnGlnGlyAsnProArgAla 190  
Db 1732 AGATCAGTTCAAGAAAGTATCCCAAGGGCC 1761  
RESULT 10  
AAF82461  
ID AAF82461 standard; cDNA; 2521 BP.  
XX  
AC AAF82461;

XX 29-JUN-2001 (first entry)  
XX Alternatively spliced human CASB6411 cDNA encoding truncated protein.  
DE  
XX  
XX Human; CASB6411; vaccine; gene therapy; immunophylaxis;  
KM ovarian cancer; colon cancer; autoimmune disease; isoform;  
KW alternative splicing; ss.  
XX  
XX Homo sapiens.  
FH Key Location/Qualifiers  
FT CDS 382..846  
FT /\*tag= a  
FT /product= "truncated CASB6411"  
XX  
XX WO200123417-A2.  
XX  
XX 05-APR-2001.  
XX  
XX 27-SEP-2000; 2000WO-EP09500.  
XX  
XX 30-SEP-1999; 99GB-0023154.  
PR 07-JUL-2000; 2000GB-0016839.  
XX  
XX (SMIK ) SMITHKLINE BEECHAM BIOLOGICALS.  
XX  
XX Vinals De Bassols YC;  
PI  
XX WPI; 2001-316133/33.  
DR P-PsDB; AAB83080.  
XX  
XX Novel CASB6411 polypeptides useful in diagnostics, and as vaccines for  
PT prophylactic and therapeutic treatment of cancers, particularly ovarian  
PT and colon cancers, autoimmune diseases and related conditions -  
XX  
XX Claim 11; Page 64-65; 95pp; English.  
XX  
XX The present sequence encodes a truncated CASB6411 polypeptide. It  
CC is generated by alternative splicing of the full length human cDNA  
CC sequence of CASB6411. The invention relates to CASB6411 polypeptides  
CC comprising a sequence having at least 70% identity to a sequence of  
CC 460 or 154 amino acids fully defined in the specification. CASB6411  
CC polypeptides and polynucleotides are useful for treating a subject by  
CC immunophylaxis or therapy. The CASB6411 polypeptides are useful in  
CC diagnostics, and as vaccines for prophylactic and therapeutic treatment  
CC of cancers, particularly ovarian and colon cancers, autoimmune diseases  
CC and related conditions. CASB6411 polypeptides are also useful for the  
CC structure-based design of agonists, antagonists or inhibitors of the  
CC polypeptide.  
XX  
XX Sequence 2521 BP; 662 A; 583 C; 583 G; 693 T; 0 other;  
SQ  
Alignment Scores:  
Pred. No.: 4.76e-183 Length: 2521  
Score: 190.00 Matches: 190  
Percent Similarity: 100.00% Conservative: 0  
Best Local Similarity: 100.00% Mismatches: 0  
Query Match: 100.00% Indels: 0  
DB: 22 Gaps: 0  
US-09-895-298A-83 (1-190) x AAF82461 (1-2521)  
QY 1 MetMetAsnPhgInProProSerIysAlaTrpArgAlaSerGlnMetMetThrPhePhe 20  
DB 1306 ATGATGAAATTTCAGCCCTCCGAGCAAGCCCTGGCGGCTCAGAGATGAGACTTCTTC 1365  
QY 21 IlePheLeuLeuPhePheProSerPheThrGlyValLeuCysThrLeuAlaIleThrIle 40  
DB 1366 ATCTCTTGCTCTTTTCCATCCTTCACCGGGGTCTGTGCACCCCTGCATCACCATC 1425  
QY 41 TrpArgLeuLysProSerAlaAspCysGlyProPheArgGlyLeuProLeuPheIleHis 60  
|||||

DB 1426 TGGAGATTGAAGCCTTCAGCTGACTGTGGCCCTTTTCGAGGTCTGCTCTTCATTTCAC 1485  
QY 61 SerIleTyrSerTrpIleAspThrLeuSerThrArgProGlyTyrLeuTrpValIleTip 80  
DB 1486 TCCATCTACAGCTGGATGAGCACCCTTAAGTACACGCGCTGCTTACCTGTGGTGGTGG 1545  
QY 81 IleTyrArgAsnLeuIleGlySerValHisPhePheIleLeuThrLeuIleValLeu 100  
DB 1546 ATCTATCGGAACCTCATTTGGAAGTGTGCACCTTCTTTTCATCTCACCCTCATTTGTCTA 1605  
QY 101 IleIleThrTyrLeuTyrTrpGlnIleThrGluGlyArgLysIleMetIleArgLeuLeu 120  
DB 1606 ATCATCACCCTATCTTACTGCGACATCACAGAGGAAGGAATATGATTAAGGCTGCTC 1665  
QY 121 HisGluGlnIleIleAsnGluGlyLysAspLysMetPheLeuIleGluLysLeuIleLys 140  
DB 1666 CATGAGCAGATCATTTAATGAGGGCAAGATAAATGTCTGTAGAAATAATTGATCAG 1725  
QY 141 LeuGlnAspMetGluLysLysAlaAsnProSerSerLeuValLeuGluArgGluVal 160  
DB 1726 CTGCAGGATATGAGAGAAGCAAAACCCAGCTCACTTGTCTGGAAGAGAGAGAGTGTG 1785  
QY 161 GluGlnGlnGlyPheLeuHisLeuGlyGluHisAspGlySerLeuAspLeuArgSerArg 180  
DB 1786 GAGCAACAAGGCTTTTGTGATTTGGGGGAACATGATGCGAGCTTGGACTTGGCATCTAGA 1845  
QY 181 ArgSerValGlnGluGlyAsnProArgAla 190  
DB 1846 AGATCAGTTCAGAGAAGGTAAATCCCAAGGCC 1875  
RESULT 11  
ABV22463  
ID ABV22463 standard; cDNA; 1194 BP.  
XX AC ABV22463;  
XX  
XX 13-SEP-2002 (first entry)  
XX  
XX Human prostate expression marker cDNA 22454.  
DE  
XX  
XX Human; prostate cancer; cytostatic; carcinogen; pharmacodynamic marker;  
KW pharmacogenomic marker; gene; ss.  
XX  
XX Homo sapiens.  
OS  
XX WO200160860-A2.  
PN  
XX 23-AUG-2001.  
PD  
XX 20-FEB-2001; 2001WO-US05171.  
PF  
XX 17-FEB-2000; 2000US-183319P.  
PR 16-MAR-2000; 2000US-189862P.  
PR 25-MAY-2000; 2000US-207454P.  
PR 09-JUN-2000; 2000US-211314P.  
PR 18-JUL-2000; 2000US-219007P.  
PR 13-DEC-2000; 2000US-255281P.  
XX  
PA (MILL-) MILLENNIUM PREDICTIVE MEDICINE INC.  
XX  
XX Schlegel R, Endege WO, Monahan JE;  
PI  
XX  
XX WPI; 2001-662795/76.  
DR  
XX Novel isolated nucleic acid molecule associated with cancerous state of  
PT prostate cells and correlating with presence of prostate cancer, useful  
PT for detecting presence of prostate cancer, stage of prostate cancer -  
XX  
PS Claim 1; Page 3912; 11750pp; English.  
XX  
CC The invention relates to an isolated nucleic acid molecule (1) comprising  
CC a nucleotide sequence given in Tables 1-9 (ABV00010-ABV62213) of the  
CC specification or its complement. (1) is useful for:

CC (a) assessing whether a patient is afflicted with prostate cancer;  
CC (b) monitoring the progression of prostate cancer in a patient;  
CC (c) assessing the efficacy of a test compound to inhibit prostate cancer in a patient;  
CC (d) assessing the efficacy of a therapy for inhibiting prostate cancer in a patient;  
CC (e) selecting a composition for inhibiting prostate cancer in a patient;  
CC (f) assessing the prostate cell carcinogenic potential of a compound;  
CC (g) determining whether prostate cancer has metastasized in a patient;  
CC (h) assessing the aggressiveness or indolence of prostate cancer in a patient;  
CC (i) is also useful as a pharmacodynamic or pharmacogenomic marker.  
SQ Sequence 1194 BP; 288 A; 287 C; 278 G; 339 T; 2 other;  
  
Alignment Scores:  
Pred. No.: 1.05e-142 Length: 1194  
Score: 150.00 Matches: 150  
Percent Similarity: 100.00% Conservative: 0  
Best Local Similarity: 100.00% Mismatches: 0  
Query Match: 78.95% Indels: 0  
DB: 23 Gaps: 0  
  
US-09-895-298A-83 (1-190) x ABV22463 (1-1194)  
QY 1 MetMetAsnPhgInProProSerLysAlaTrpArgAlaSerGlnMetMetThrpPhe 20  
Db 531 ATGATGAATTTCCAGCCTCCGAGCAAGCCTGGCGGCTCAGATGATGACTTCTTC 590  
QY 21 IlePheLeuLeuPhePheProSerPheThrGlyValLeuGlyThrLeuAlaIleThrIle 40  
Db 591 ATCTTCTGCTCTTTTCCATCCTTCACCGGGGCTTGTGCACCTGGCCATCACCATC 650  
QY 41 TrpArgLeuLysProSerAlaAspCysGlyProPheArgGlyLeuProLeuPheIleHis 60  
Db 651 TGGAGATGAAGCCTTCAGCTGACTGTGGCCCTTTTCGAGGTCTGCTCTTCATTGAC 710  
QY 61 SerIleTyrSerTrpIleAspThrLeuSerThrArgProGlyTyrLeuTrpValValTrp 80  
Db 711 TCCATCTACAGCTGGATGACACCCCTAAGTACACGGCCTGCTACTGTGGTGTGG 770  
QY 81 IleTyrArgAsnLeuIleGlySerValHisPhePheIleLeuThrLeuIleValLeu 100  
Db 771 ATCTATCGGAACCTTCATTGGAAGTGTGACTCTTTTCATCTCACCCTCATTTGTGCTA 830  
QY 101 IleIleThrTyrLeuTyrTrpGlnIleThrGlyArgLysIleMetIleArgLeuLeu 120  
Db 831 ATCATCACCCTATCTTACTGCGACATCAGAGGGAAGAAATGATATAGAGCTGCTC 890  
QY 121 HisGlnGlnIleIleAsnGlnGlyLysAspLysMetPheLeuIleGlnLysLeuIleLys 140  
Db 891 CATGAGCAGATCATTAATGAGGGCAAGATAAATGTTCTGATGAAAAAATTGATCAAG 950  
QY 141 LeuGlnAspMetGlnLysLysAlaAsnPro 150  
Db 951 CTGCAGATATGAGAAAGCAACCA 980  
  
RESULT 12  
ABV25683  
ID ABV25683 standard; cDNA; 1194 BP.  
XX ABV25683;  
AC  
XX  
XX 16-SEP-2002 (first entry)  
DT  
XX  
XX Human prostate expression marker cDNA 25674.  
DE  
XX  
XX Human; prostate cancer; cytostatic; carcinogen; pharmacodynamic marker;  
KW pharmacogenomic marker; gene; ss.  
XX  
OS Homo sapiens.  
XX  
PN WO200160860-A2.

XX  
PD 23-AUG-2001.  
XX  
XX 20-FEB-2001; 2001WO-US05171.  
PF  
XX  
XX 17-FEB-2000; 2000US-183319P.  
PR 16-MAR-2000; 2000US-189862P.  
PR 25-MAY-2000; 2000US-207454P.  
PR 09-JUN-2000; 2000US-211314P.  
PR 18-JUL-2000; 2000US-219007P.  
PR 13-DEC-2000; 2000US-255281P.  
XX  
XX (MILL-) MILLENNIUM PREDICTIVE MEDICINE INC.  
PI Schlegel R, Endege WO, Monahan JE;  
XX  
XX WPI; 2001-662795/76.  
DR  
XX  
XX Novel isolated nucleic acid molecule associated with cancerous state of  
PT prostate cells and correlating with presence of prostate cancer, useful  
PT for detecting presence of prostate cancer, stage of prostate cancer -  
XX  
XX Claim 1; Page 5146-5147; 11750pp; English.  
PS  
XX  
XX The invention relates to an isolated nucleic acid molecule (I) comprising  
CC a nucleotide sequence given in Tables 1-9 (ABV00010-ABV62213) of the  
CC specification or its complement. (I) is useful for:  
CC (a) assessing whether a patient is afflicted with prostate cancer;  
CC (b) monitoring the progression of prostate cancer in a patient;  
CC (c) assessing the efficacy of a test compound to inhibit prostate cancer in a patient;  
CC (d) assessing the efficacy of a therapy for inhibiting prostate cancer in a patient;  
CC (e) selecting a composition for inhibiting prostate cancer in a patient;  
CC (f) assessing the prostate cell carcinogenic potential of a compound;  
CC (g) determining whether prostate cancer has metastasized in a patient;  
CC (h) assessing the aggressiveness or indolence of prostate cancer in a patient;  
CC (i) is also useful as a pharmacodynamic or pharmacogenomic marker.  
SQ Sequence 1194 BP; 288 A; 287 C; 278 G; 339 T; 2 other;  
  
Alignment Scores:  
Pred. No.: 1.05e-142 Length: 1194  
Score: 150.00 Matches: 150  
Percent Similarity: 100.00% Conservative: 0  
Best Local Similarity: 100.00% Mismatches: 0  
Query Match: 78.95% Indels: 0  
DB: 23 Gaps: 0  
  
US-09-895-298A-83 (1-190) x ABV25683 (1-1194)  
QY 1 MetMetAsnPhgInProProSerLysAlaTrpArgAlaSerGlnMetMetThrpPhe 20  
Db 531 ATGATGAATTTCCAGCCTCCGAGCAAGCCTGGCGGCTCAGATGATGACTTCTTC 590  
QY 21 IlePheLeuLeuPhePheProSerPheThrGlyValLeuGlyThrLeuAlaIleThrIle 40  
Db 591 ATCTTCTGCTCTTTTCCATCCTTCACCGGGGCTTGTGCACCTGGCCATCACCATC 650  
QY 41 TrpArgLeuLysProSerAlaAspCysGlyProPheArgGlyLeuProLeuPheIleHis 60  
Db 651 TGGAGATGAAGCCTTCAGCTGACTGTGGCCCTTTTCGAGGTCTGCTCTTCATTGAC 710  
QY 61 SerIleTyrSerTrpIleAspThrLeuSerThrArgProGlyTyrLeuTrpValValTrp 80  
Db 711 TCCATCTACAGCTGGATGACACCCCTAAGTACACGGCCTGCTACTGTGGTGTGG 770  
QY 81 IleTyrArgAsnLeuIleGlySerValHisPhePheIleLeuThrLeuIleValLeu 100  
Db 771 ATCTATCGGAACCTTCATTGGAAGTGTGACTCTTTTCATCTCACCCTCATTTGTGCTA 830  
QY 101 IleIleThrTyrLeuTyrTrpGlnIleThrGlyArgLysIleMetIleArgLeuLeu 120

Db	831	ATCATCACCTATCTTACTTGCGAGATCACAGAGGAAGGATTTATGATAAGGCTGCTC	890
Qy	121	HisGluglnIleIleasnGlucIylLysAspLysMetPheLeuIleGlulysLeuIleLys	140
Db	891	CATGAGCAGATCATTTAATGAGGGCAAAAGATAAAATGTTCTGATAGAAAAAATTGATCAAG	950
Qy	141	LeuGlnAspMetGlulysLysAlaAsnPro	150
Db	951	CTGCGAGATATGAGAGAAAGCAACCA	980
RESULT 13			
ABV282278	ID	ABV282278 standard; cDNA; 1194 BP.	
XX	AC	ABV282278;	
XX	DT	16-SEP-2002 (first entry)	
XX	DE	Human prostate expression marker CDNA 28269.	
XX	KW	Human; prostate cancer; cytostatic; carcinogen; pharmacodynamic marker;	
XX	KW	pharmacogenomic marker; gene; ss.	
XX	OS	Homo sapiens.	
XX	PN	WO200160860-A2.	
XX	PD	23-AUG-2001.	
XX	PF	20-FEB-2001; 2001WO-US05171.	
XX	PR	17-FEB-2000; 2000US-183319P.	
XX	PR	16-MAR-2000; 2000US-189862P.	
XX	PR	25-MAY-2000; 2000US-207454P.	
XX	PR	09-JUN-2000; 2000US-211314P.	
XX	PR	18-JUL-2000; 2000US-219007P.	
XX	PR	13-DEC-2000; 2000US-255281P.	
XX	PA	(MILL-) MILLENNIUM PREDICTIVE MEDICINE INC.	
XX	PI	Schlegel R, Endege WO, Monahan JE;	
XX	DR	WPI; 2001-662795/76.	
XX	PT	Novel isolated nucleic acid molecule associated with cancerous state of prostate cells and correlating with presence of prostate cancer, useful for detecting presence of prostate cancer, stage of prostate cancer -	
XX	PS	Claim 1; Page 5881-5882; 11750pp; English.	
XX	CC	The invention relates to an isolated nucleic acid molecule (I) comprising a nucleotide sequence given in Tables 1-9 (ABV00010-ABV62213) of the specification or its complement. (I) is useful for:	
XX	CC	(a) assessing whether a patient is afflicted with prostate cancer;	
XX	CC	(b) monitoring the progression of prostate cancer in a patient;	
XX	CC	(c) assessing the efficacy of a test compound to inhibit prostate cancer in a patient;	
XX	CC	(d) assessing the efficacy of a therapy for inhibiting prostate cancer in a patient;	
XX	CC	(e) selecting a composition for inhibiting prostate cancer in a patient;	
XX	CC	(f) assessing the prostate cell carcinogenic potential of a compound;	
XX	CC	(g) determining whether prostate cancer has metastasized in a patient;	
XX	CC	(h) assessing the aggressiveness or indolence of prostate cancer in a patient;	
XX	CC	(I) is also useful as a pharmacodynamic or pharmacogenomic marker.	
XX	SQ	Sequence 1194 BP; 288 A; 287 C; 278 G; 339 T; 2 other;	
Alignment Scores:			
Pred. No.:	1.05e-142	Length:	1194
Score:	150.00	Matches:	150
Percent Similarity:	100.00%	Conservative:	0

Best Local Similarity:	100.00%	Mismatches:	0
Query Match:	78.95%	Indels:	0
DB:	23	Gaps:	0
US-09-895-298A-83 (1-190) x ABV28278 (1-1194)			
QY	1 MetMetAsnPhcInProProSerLysAlaTrpArgAlaSerGlnMetMetTrpPhe	20	
DB	531 ATGATGAATTTCCAGCCTCCGAGCAAGAGCTGGCGGCTCACAGATGATGACTTCTTC	590	
QY	21 IlePheLeuLeuPhePheProSerPheThrGlyValLeuCysThrLeuAlaIleThrIle	40	
DB	591 ATCTTCTTGCTCTTTTCCCATCTTCACCGGGGCTTGTCACCCCTGGCCATCACCATC	650	
QY	41 TrpArgLeuLysProSerAlaAspCysGlyProPheArgGlyLeuProLeuPheIleHis	60	
DB	651 TGGAGATTGAAGCCTTCAGCTGACTGTGGCCCTTTTCGAGGCTGTGCTCTCTCATTTAC	710	
QY	61 SerIleYrSerTrpIleAspThrLeuSerThrArpProGlyTrpLeuTrpValValTrp	80	
DB	711 TCCATCTACAGCTGATCGACACCCCTAAGTACAGCGCTGCTACCTGTGGTGTGTGG	770	
QY	81 IleTyraArgAsnLeuIleGlySerValHisPhePheIleLeuThrLeuIleValLeu	100	
DB	771 ATCTATCGGAACCTCATTTGGAAGTGTGCACCTTTTTCATCTCACCCTCATTTGTGCTA	830	
QY	101 IleIleThrTrpLeuIleTrpGlnIleThrGluGlyArgLysIleMetIleArgLeuLeu	120	
DB	831 ATCATCACCTATCTTTACTGGCAGATCACAGAGGAAGAGATTATATTAAGCTGTCTC	890	
QY	121 HisGluGlnIleIleAsnGluGlyLysAspLysMetPheLeuIleGluLysLeuIleLys	140	
DB	891 CATGAGCAGATCATTAATGAGGGCAAGATAAAATGTTCCGTAGTAAAGAAATTGATCAAG	950	
QY	141 LeuGlnAspMetGluLysLysAlaAsnPro	150	
DB	951 CTGCAGGATATGAGAAAGCAAAACCA	980	
RESULT 14			
ID	AA118591 standard; cDNA; 470 BP.		
XX	AA118591;		
AC			
XX			
DT	07-DEC-2001 (first entry)		
XX			
DE	Human breast cancer expressed polynucleotide 11048.		
XX			
KW	Human; breast cancer; cell marker; cytosolic; ss.		
XX			
OS	Homo sapiens.		
XX			
XX			
PN	WO200151628-A2.		
XX			
PD	19-JUL-2001.		
XX			
PF	10-JAN-2001; 2001WO-US00798.		
XX			
PR	14-JAN-2000; 2000US-0176077.		
XX			
PR	14-MAR-2000; 2000US-0189167.		
XX			
PR	24-MAR-2000; 2000US-0192099.		
XX			
PR	29-MAR-2000; 2000US-0193480.		
XX			
PR	15-MAY-2000; 2000US-0205230.		
XX			
PR	09-JUN-2000; 2000US-0211315.		
XX			
PR	25-JUL-2000; 2000US-0220534.		
XX			
PA	(MILL-) MILLENNIUM PREDICTIVE MEDICINE INC.		
XX			
PI	Lillie J, Xu Y, Wang Y, Steinmann K;		
XX			
DR	WPI; 2001-451856/48.		
XX			
PT	New peptide useful as a marker for the diagnosis of breast cancer		

XX Claim 1; Page 1968; 3695pp; English.  
PS  
XX  
CC The invention relates to human breast cancer expressed polynucleotides  
CC (AAL07544-AAL26789) and methods of assessing whether a patient is  
CC afflicted with breast cancer by examining the correlation between the  
CC expression of certain markers and the cancerous state of breast cells.  
CC The polynucleotides and encoded polypeptides are potential markers for  
CC detecting, diagnosing, monitoring, characterizing treating and  
CC potentially preventing breast cancer. The polynucleotides and encoded  
CC polypeptides are also useful for isolating compounds with cytostatic  
CC activity.  
XX  
SQ Sequence 470 BP; 144 A; 92 C; 116 G; 118 T; 0 other;  
XX  
Alignment Scores:  
Pred. No.: 1.32e-112 Length: 470  
Score: 120.00 Matches: 120  
Percent Similarity: 100.00% Conservative: 0  
Best Local Similarity: 100.00% Mismatches: 0  
Query Match: 63.16% Indels: 0  
DB: 22 Gaps: 0  
US-09-895-298a-83 (1-190) x AAL18591 (1-470)  
QY 71 ThrArgProGlyTyrLeuTrpValValTrpIleTyrArgAsnLeuIleGlySerValHis 90  
Db 5 ACACGGCCCTGGCTACCTGTGGTGTGGATCATCGAACCCTCATTTGAGAGTGTGCAC 64  
QY 91 PhePhePheIleLeuThrLeuIleValLeuIleIleThrTyrLeuTyrTrpGlnIleThr 110  
Db 65 TTCTTTTTCATCCTCAACCCTCATTTGCTAATCATCACCTATCTTAACTGGCAGATCACA 124  
QY 111 GluGlyArgLysIleMetIleArgLeuLeuHisGluGlnIleIleAsnGluGlyLysAsp 130  
Db 125 GAGGGAAGAGATATGATAGAGCTGCTCCATGACAGACATCATTAATGAGGCAAGAT 184  
QY 131 LysMetPheLeuIleGluLysLeuIleLysLeuGlnAspMetGluLysLysAlaAsnPro 150  
Db 185 AAAATGTCCTGATGAAAAAATTTGATCAAGCTCGAGATATGAGAAAGAAACCAACCC 244  
QY 151 SerSerLeuValLeuGluArgArgGluValGluGlnGlnGlyPheLeuHisLeuGlyGlu 170  
Db 245 AGCTCAGTTGTTCTGGAAGAGAGAGAGGTGAGACCAACAGGCTTTTTCATTGGGGGAA 304  
QY 171 HisAspGlySerLeuAspLeuArgSerArgSerValGlnGlnGlyAsnProArgAla 190  
Db 305 CATGATGGCAGTCTTGACTCGGATCTAGAGATCAAGTCAAGAAGCTAATCCAAAGGGCC 364  
RESULT 15  
AAL09919  
ID AAL09919 standard; cDNA; 501 BP.  
XX  
AC AAL09919;  
XX  
DT 07-DEC-2001 (first entry)  
XX  
DE Human breast cancer expressed polynucleotide 2376.  
XX  
KW Human; breast cancer; cell marker; cytostatic; ss.  
XX  
OS Homo sapiens.  
XX  
PN WO200151628-A2.  
XX  
PD 19-JUL-2001.  
XX  
PF 10-JAN-2001; 2001WO-US00798.  
XX  
PR 14-JAN-2000; 2000US-0176077.  
PR 14-MAR-2000; 2000US-0189167.  
PR 24-MAR-2000; 2000US-0192099.  
PR 29-MAR-2000; 2000US-0193480.

PR 15-MAY-2000; 2000US-0205230.  
PR 09-JUN-2000; 2000US-0211315.  
PR 25-JUL-2000; 2000US-0220534.  
XX  
PA (MILL-) MILLENNIUM PREDICTIVE MEDICINE INC.  
PI Lillie J, Xu Y, Wang Y, Steinmann K;  
XX  
DR WPI; 2001-451856/48.  
XX  
PT New peptide useful as a marker for the diagnosis of breast cancer -  
XX  
PS Claim 1; Page 455; 3695pp; English.  
XX  
SQ Sequence 501 BP; 147 A; 101 C; 128 G; 122 T; 3 other;  
XX  
Alignment Scores:  
Pred. No.: 2.04e-102 Length: 501  
Score: 110.00 Matches: 110  
Percent Similarity: 100.00% Conservative: 0  
Best Local Similarity: 100.00% Mismatches: 0  
Query Match: 57.89% Indels: 0  
DB: 22 Gaps: 0  
US-09-895-298a-83 (1-190) x AAL09919 (1-501)  
QY 71 ThrArgProGlyTyrLeuTrpValValTrpIleTyrArgAsnLeuIleGlySerValHis 90  
Db 36 ACACGGCCCTGGCTACCTGTGGTGTGGATCATCGAACCCTCATTTGAGAGTGTGCAC 95  
QY 91 PhePhePheIleLeuThrLeuIleValLeuIleIleThrTyrLeuTyrTrpGlnIleThr 110  
Db 96 TTCTTTTTCATCCTCAACCCTCATTTGCTAATCATCACCTATCTTAACTGGCAGATCACA 155  
QY 111 GluGlyArgLysIleMetIleArgLeuLeuHisGluGlnIleIleAsnGluGlyLysAsp 130  
Db 156 GAGGGAAGAGATATGATAGAGCTGCTCCATGACAGACATCATTAATGAGGCAAGAT 215  
QY 131 LysMetPheLeuIleGluLysLeuIleLysLeuGlnAspMetGluLysLysAlaAsnPro 150  
Db 216 AAAATGTCCTGATGAAAAAATTTGATCAAGCTCGAGATATGAGAAAGAAACCAACCC 275  
QY 151 SerSerLeuValLeuGluArgArgGluValGluGlnGlnGlyPheLeuHisLeuGlyGlu 170  
Db 276 AGCTCAGTTGTTCTGGAAGAGAGAGAGGTGAGACCAACAGGCTTTTTCATTGGGGGAA 335  
QY 171 HisAspGlySerLeuAspLeuArgSerArg 180  
Db 336 CATGATGGCAGTCTTGACTTGGATCTAGA 365

Search completed: November 9, 2002, 07:45:52  
Job time : 305 secs